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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/501,284	02/07/2005	Gesine Schliecker	I-2002.001 US	5686	
	7590 04/08/201 ng-Plough Animal Hea	EXAMINER			
Patent Dept. K-6-1, 1990 2000 Galloping Hill Road Kenilworth, NJ 07033-0530			PERREIRA, MELISSA JEAN		
			ART UNIT	PAPER NUMBER	
			1618		
			NOTIFICATION DATE	DELIVERY MODE	
			04/08/2010	ELECTRONIC	

## Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

patents@spcorp.com jill.corcoran@spcorp.com

## Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)	
10/501,284	SCHLIECKER ET AL.	
Examiner	Art Unit	

	MELISSA PERREIRA	1618				
The MAILING DATE of this communication appea	ars on the cover sheet with the c	correspondence add	ress			
THE REPLY FILED <u>19 March 2010</u> FAILS TO PLACE THIS API	PLICATION IN CONDITION FOR A	ALLOWANCE.				
1.  The reply was filed after a final rejection, but prior to or on application, applicant must timely file one of the following rapplication in condition for allowance; (2) a Notice of Appe for Continued Examination (RCE) in compliance with 37 C periods:	eplies: (1) an amendment, affidavit al (with appeal fee) in compliance	t, or other evidence, w with 37 CFR 41.31; or	hich places the (3) a Request			
a) The period for reply expiresmonths from the mailing	date of the final rejection.					
b) The period for reply expires on: (1) the mailing date of this Ac no event, however, will the statutory period for reply expire la	ter than SIX MONTHS from the mailing	g date of the final rejection	n.			
Examiner Note: If box 1 is checked, check either box (a) or (b) MONTHS OF THE FINAL REJECTION. See MPEP 706.07(f)						
Extensions of time may be obtained under 37 CFR 1.136(a). The date of have been filed is the date for purposes of determining the period of extender 37 CFR 1.17(a) is calculated from: (1) the expiration date of the state forth in (b) above, if checked. Any reply received by the Office later that may reduce any earned patent term adjustment. See 37 CFR 1.704(b). NOTICE OF APPEAL	ension and the corresponding amount of cortened statutory period for reply original cortened statutory period	of the fee. The appropria nally set in the final Offic	ate extension fee e action; or (2) as			
2. The Notice of Appeal was filed on A brief in compl	ance with 37 CFR 41.37 must be f	iled within two month	s of the date of			
filing the Notice of Appeal (37 CFR 41.37(a)), or any exten Notice of Appeal has been filed, any reply must be filed with AMENDMENTS	sion thereof (37 CFR 41.37(e)), to	avoid dismissal of the				
 3. ☐ The proposed amendment(s) filed after a final rejection, b	ut prior to the date of filing a brief,	will not be entered be	cause			
(a) They raise new issues that would require further con	· · · · · · · · · · · · · · · · · · ·					
(b) ☐ They raise the issue of new matter (see NOTE below	•					
(c) They are not deemed to place the application in bette	er form for appeal by materially rec	ducing or simplifying t	ne issues for			
appeal; and/or (d) ☐ They present additional claims without canceling a c	orresponding number of finally reje	ected claims				
NOTE: (See 37 CFR 1.116 and 41.33(a)).	orresponding number of initially reje	otod oldiirio.				
4. The amendments are not in compliance with 37 CFR 1.12	1. See attached Notice of Non-Co	mpliant Amendment (	PTOL-324).			
5. Applicant's reply has overcome the following rejection(s):			,			
6. Newly proposed or amended claim(s) would be allow	owable if submitted in a separate, t	imely filed amendmer	nt canceling the			
non-allowable claim(s).  7. For purposes of appeal, the proposed amendment(s): a)	will not be entered or b) \( \square\) will	l he entered and an e	volenation of			
how the new or amended claims would be rejected is provi The status of the claim(s) is (or will be) as follows:		i pe entereu and an e.	хріапацоп оі			
Claim(s) allowed:						
Claim(s) objected to: Claim(s) rejected:						
Claim(s) withdrawn from consideration:						
AFFIDAVIT OR OTHER EVIDENCE						
<ol> <li>The affidavit or other evidence filed after a final action, but because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).</li> </ol>						
<ol> <li>The affidavit or other evidence filed after the date of filing a entered because the affidavit or other evidence failed to over the entered because the affidavit or other evidence.</li> </ol>	ercome <u>all</u> rejections under appea	ıl and/or appellant fail	s to provide a			
showing a good and sufficient reasons why it is necessary and was not earlier presented. See 37 CFR 41.33(d)(1).  0. The affidavit or other evidence is entered. An explanation of the status of the claims after entry is below or attached.						
REQUEST FOR RECONSIDERATION/OTHER		ing to botom or analytic	<b>-</b>			
11.   ☐ The request for reconsideration has been considered but see below.	,	condition for allowan	ce because:			
<ul><li>12. ☐ Note the attached Information <i>Disclosure Statement</i>(s). (I</li><li>13. ☐ Other:</li></ul>	PTO/SB/08) Paper No(s)					
/Michael G. Hartley/	/Melissa Perreira/					
Supervisory Patent Examiner, Art Unit 1618	Examiner, Art Unit 1618					
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Claims 1-19 and 21-26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Krone et al. (US 5,391,696) in view of Lewis (US 5,838,571) and in further view of Suzuki et al. (US 6,015,789) and Remington's Pharmaceutical Sciences 1990 18th Ed. Chpt. 89.

Applicant asserts that the references cannot render obvious amended claims 1,14,24 and 26, as neither references themselves nor the inferences and creative steps that a person of ordinary skill in the art would have employed at the time of the invention taught or suggested a polytartrate composition having "a lag phase of a predetermined time" in the release of a pharmaceutical composition as recited by amended claims 1,24,26, or actually "determining a time length of the lag phase" as recited by amended claim 14.

Krone et al. teaches polytartrate formulations which may comprise tablets formed via compaction/compression and do not comprise a barrier structure. Krone et al. teaches that polytartrate preparations have a decreased "initial burst" which implies that they have a second burst/release. In regards to the "lag time" the specification recites (p13, lines 17-20 and 30-33), "a secondary "lag phase" of low or no release of the drug followed by a second burst". Therefore, the polytartrate tablet formulations which are prepared via compaction/compression of Krone et al. encompass the composition of the instant claims as they have a first "initial burst", an implied second burst and the phase between the bursts (i.e. lag phase) which may release drug. Therefore, the time between bursts of the polytartrate formulations of Krone et al. encompass the "lag phase" of the instant claims.

Also, the recitation, "determining a time of the lag phase", is a mental step and does not contain any active technique (manual steps) for determining a time of the lag phase. There are no specific active steps to define how such determining is performed or which limit the amount of time of the lag phase.

Applicant asserts that Krone et al. teaches that in general after a considerable "initial burst" only a small to moderate release rate is effected. Thus, Krone et al. teaches that there can be an "initial burst" followed by a steady state rate without the need for any subsequent burst, that nothing in the cited references implies that an "initial burst" must be followed by a second burst as would be understood by one of ordinary skill in the art and that the "initial burst" simply refers to the immediate release of the available pharmaceutical from the composition, which is then followed by a more sustained slow or moderate release rate.

Krone et al. teaches that in general after a considerable "initial burst" only a small to moderate release rate is effected with regards to the prior art polyester degradable polymer pharmaceutical systems (Krone et al. column 1, lines 41-42).

Krone et al. teaches polytartrate formulations which may comprise tablets formed via compaction/compression and do not comprise a barrier structure. Krone et al. teaches that polytartrate preparations have a decreased "initial burst" which implies that they have a second burst/release. In regards to the "second burst", the specification recites that the "second booster dose (burst)" (p6, lines 1-7; p13, lines 14-16) occurs over a period of, preferably, 1-4 days while the "initial burst" occurs over 1-3 days.

According to the specification, a burst occurs over a period of days and therefore the steady state release rate/moderate release, as asserted by the applicant, of the "second burst" of Krone et al. encompasses the "second burst" of the instant invention.

The instant claims 1-18,21,22 and 24-26 do not recite a "second burst".

Applicant asserts that Krone et al. teaches compositions displaying "a strongly decreased 'initial burst' when they were used for depot preparations of pharmaceuticals" and thus teaches away from the claimed compositions as the "initial burst" is unwated.

Krone et al. teaches that the initial active substance release (initial burst) over 24 hours may be 55% (column 15, table 1 and lines 29-39).

The specification recites that the "initial burst" occurs over 1-3 days (p12, lines 22-24) and therefore the initial burst of Krone et al. (55% over 24 hours) encompasses the initial burst of the instant invention (over a period of 1-3 days).